

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re: Williams *et al.*

Confirmation No.: 1920

Serial No.: 10/662,757

Group Art Unit: 1792

Filed: September 15, 2003

Examiner: James Lin

For: *INTRALUMINAL PROSTHESES AND CARBON DIOXIDE-ASSISTED
METHODS OF IMPREGNATING SAME WITH PHARMACOLOGICAL AGENTS*

Date: October 1, 2008

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APPELLANT'S SUPPLEMENTAL BRIEF ON APPEAL UNDER 37 C.F.R. § 41.37

Sir:

This Supplemental Appeal Brief is filed pursuant to the "Notice of Non-Compliant Appeal Brief" mailed September 5, 2008 and the "Notice of Appeal to the Board of Patent Appeals and Interferences" filed on May 6, 2008. Appellants note that this Supplemental Appeal Brief incorporates amendments to the Brief provided on July 16, 2008, which were provided in response to a Notice of Non-Compliant Appeal Brief mailed July 11, 2008.

Appellant notes that the Appeal fee was paid with the Appeal Brief of June 30, 2008. Accordingly, **no additional fee is believed due**. However, any additional fees believed to be due in connection with this paper may be charged to our Deposit Account No. 50-0220

REAL PARTY IN INTEREST

The real party in interest is Synecor, LLC, the assignee of the rights to this application by virtue of assignment from the inventors to Synecor, LLC, recorded at the United States Patent and Trademark Office on December 17, 2003 on Reel 014795, Frame 0446.

RELATED APPEALS AND INTERFERENCES

Appellants are aware of no related appeals or interferences that would be affected by the present appeal.

STATUS OF CLAIMS

Claims 73-104 are pending in the present application as of the filing date of this Appeal Brief. Claims 1-72 have been cancelled from the present application. As of the filing date of this Appeal Brief, Claims 73-104 stand finally rejected under 35 U.S.C. § 103(a) as noted in the Final Office Action mailed February 12, 2008 and the Advisory Action mailed April 25, 2008.

Appellants appeal the rejection of Claims 73-104. A copy of Claims 73-104 is attached hereto as **Claims Appendix**, presenting the claims at issue as twice rejected in the Final Office Action dated February 28, 2007 and the Advisory Action mailed April 25, 2008.

STATUS OF AMENDMENTS

All amendments made by Appellants during prosecution are believed to be entered as indicated by the Final Office Action dated February 12, 2008.

SUMMARY OF CLAIMED SUBJECT MATTER

The present invention is directed to a method of impregnating an intraluminal prosthesis with pharmacological agents for delivery within a body of a subject. *See*, Specification, for example, at least, on page 4, lines 1-18; and Figures 1-3. Accordingly, independent **Claim 73** is directed to a method of:

immersing an intraluminal prosthesis in a mixture of a carrier fluid and a pharmacological agent, wherein the intraluminal prosthesis comprises non-layered polymeric material (Fig.1, block 100);

pressurizing the mixture of carrier fluid (Fig.1, block 110) and pharmacological agent for a time sufficient to cause the carrier fluid and pharmacological agent to at least partially penetrate the non-layered polymeric material;

removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material (Fig.1, block 120) in a predetermined concentration gradient, wherein the concentration gradient defines an elution profile of the pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject. *See Specification*, for example, at least on page 4, lines 2-19; on page 15, lines 6-12; on page 16, line 17 through page 17, line 2; on page 18, lines 12-29; and Fig. 1.

The controlled conditions involve controlling at least one parameter in a predetermined pattern, such parameters include temperature, rate of temperature change, pressure, rate of pressure change, carrier fluid quantity, and rate of carrier fluid quantity. *Specification*, page 16, lines 22-28. The carrier fluid can be carbon dioxide. *Specification*, page 5, lines 4-25. Further, one or more portions of the intraluminal prosthesis may be masked with a protective layer of material prior to immersion in a mixture of carrier gas and pharmacological agent so as to create portions or regions of the polymeric material having different concentrations of the pharmacological agent entrapped in it, or to partition one pharmacological agent in one region of the prosthesis from another pharmacological agent in a second (or third or fourth) region of the prosthesis. *See Specification*, for example, at least on page 18, line 29 through page 19, line 1; and on page 19, lines 2-4.

Independent **Claim 88** is directed to a method of:

immersing an intraluminal prosthesis in a mixture of a carrier fluid and a pharmacological agent, wherein the intraluminal prosthesis comprises non-layered polymeric material;

placing the intraluminal prosthesis within a pressure vessel (Fig. 2, block 200);

pressurizing the interior of the pressure vessel with an inert gas to a predetermined pressure (Fig. 2, block 210), wherein the inert gas is selected from the group consisting of helium, nitrogen, and argon;

supplying a mixture of a carrier fluid and a pharmacological agent into the pressure vessel (Fig. 2, block 220);

exposing the non-layered polymeric material and the mixture of carrier fluid and pharmacological agent in the pressure vessel for a time such that the carrier fluid and pharmacological agent at least partially penetrate the non-layered polymeric material (Fig. 2, block 230); and

releasing the pressure in the pressure vessel over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient (Fig. 2, block 240), wherein the concentration gradient defines an elution profile of the pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject. *See* Specification, for example, at least on page 4, line 19 through page 5, line 2; on page 16, line 17 through page 17, line 2; on page 18, lines 12-28; on page 19, lines 9-17; and in Fig. 2.

The controlled conditions recited in Claim 88 involve controlling at least one parameter in a predetermined pattern, such parameters include temperature, rate of temperature change, pressure, rate of pressure change, carrier fluid quantity, and rate of carrier fluid quantity. *See* Specification, for example, at least on page 16, lines 22-28.

Independent **Claim 99** is directed to a method of:

masking portions of an intraluminal prosthesis with a protective layer of material such that the intraluminal prosthesis has first and second unmasked portions, wherein the intraluminal prosthesis comprises non-layered polymeric material (*See* Specification, for example, at least on page 18, line 29 through page 19, line 1);

immersing the intraluminal prosthesis in a mixture of a carrier fluid and first and second pharmacological agents (*See* Specification, for example, at least original claim 1; Fig.6; and on page 24, lines 8-28);.

pressurizing the mixture of carrier fluid and pharmacological agents for a time sufficient to cause the carrier fluid and the first pharmacological agent to at least partially penetrate the first unmasked portion and to cause the carrier fluid and the second pharmacological agent to at least

partially penetrate the second unmasked portion (*See* Specification, for example, at least on page 4, lines 9-14 and 24-32, on page 18 lines 22-29 and Fig. 2); and

removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and such that an amount of the first pharmacological agent remains elutably trapped within the first unmasked portion in a predetermined concentration gradient and an amount of the second pharmacological agent remains elutably trapped within the second unmasked portion in a predetermined concentration gradient, wherein each concentration gradient defines an elution profile of a respective pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject. *See* Specification, for example, at least on page 19, lines 9-17.

GROUND OF REJECTION TO BE REVIEWED ON APPEAL

1. Whether Claims 73, 74, 76, 80-84 and 86 are properly rejected under 35 U.S.C §103(a) as unpatentable over U.S. Patent Publication No. 2003/0104030 to Igaki et al.
2. Whether Claims 73, 74, 76-78, 80-82, 86, 88, 89, 91-93 and 98 are properly rejected under 35 U.S.C. §103(a) as unpatentable over European Patent No. EP 0405284 to Greiner.
3. Whether Claims 75, 99-101 and 104 are properly rejected under 35 U.S.C. §103(a) as unpatentable over U.S. Patent Publication No. 2003/0104030 to Igaki et al. in view of U.S. Patent No. 6,251,136 to Guruwaiya.
4. Whether Claims 75, 90 and 99-104 are properly rejected under 35 U.S.C. §103(a) as unpatentable over European Patent No. EP 0405284 to Greiner in view of U.S. Patent No. 6,251,136 to Guruwaiya.

5. Whether Claim 79 is properly rejected under 35 U.S.C. §103(a) as unpatentable over U.S. Patent Publication No. 2003/0104030 to Igaki et al. in view of U.S. Patent No. 6,670,398 to Edwards.

6. Whether Claims 79 and 95 are properly rejected under 35 U.S.C. §103(a) as unpatentable over European Patent No. EP 0405284 to Greiner in view of U.S. Patent No. 6,670,398 to Edwards.

7. Whether Claims 85 and 97 are properly rejected under 35 U.S.C. §103(a) as unpatentable over European Patent No. EP 0405284 to Greiner in view of PCT Publication WO 01/87368 to Mehta.

8. Whether Claim 87 is properly rejected under 35 U.S.C. §103(a) as unpatentable over U.S. Patent Publication No. 2003/0104030 to Igaki et al. in view of U.S. Patent No. 6,299,604 to Ragheb.

9. Whether Claim 87 is properly rejected under 35 U.S.C. §103(a) as unpatentable over European Patent No. EP 0405284 to Greiner in view of U.S. Patent No. 6,299,604 to Ragheb.

10. Whether Claims 81, 83, 84, 86, 93, 94, 96 and 98 are properly rejected under 35 U.S.C. §103(a) as unpatentable over European Patent No. EP 0405284 to Greiner in view of U.S. Patent Publication No. 2003/0104030 to Igaki et al.

ARGUMENT

I. 35 U.S.C. §103 Analysis

To establish a *prima facie* case of obviousness, the prior art reference or references when combined must teach or suggest all the recitations of the claims, and there must be some suggestion or motivation, either in the references themselves or in the knowledge generally available to one of ordinary skill in the art, to modify the reference or to combine reference

teachings. M.P.E.P. §2143. A patent composed of several elements is not proved obvious merely by demonstrating that each of its elements was, independently, known in the prior art. *KSR Int'l Co. v. Teleflex Inc.*, 550 U. S. 1, 15 (2007). A corollary principle is that, when the prior art teaches away from combining certain known elements, discovery of a successful means of combining them is more likely to be unobvious. *Id.* at 12. If a technique has been used to improve one device, and a person of ordinary skill in the art would recognize that it would improve similar devices in the same way, using the technique is obvious unless its actual application is beyond his or her skill. *Id.* at 13. A Court must ask whether the improvement is more than the predictable use of prior art elements according to their established functions. *Id.* at 13. When it is necessary for a Court to look at interrelated teachings of multiple patents, the Court must determine whether there was an apparent reason to combine the known elements in the fashion claimed by the patent at issue. *Id.* at 14.

Appellants respectfully submit that the pending independent claims are patentable over the cited references for at least the reason that the cited references do not disclose or suggest many of the recitations of the claims. The patentability of the pending claims is discussed in detail hereinafter.

II. Claims 73, 74, 76, 80-84 and 86 are patentable over U.S. Patent Publication No. 2003/0104030 to Igaki et al.

Claims 73, 74, 76, 80-84 and 86 stand rejected under 35 U.S.C. § 103(a) as allegedly obvious over U.S. Patent Publication No. 2003/0104030 to Igaki et al. (hereinafter "Igaki"). Appellants respectfully disagree and request reversal of this rejection.

A. Independent Claim 73.

Appellants' independent Claim 73 recites a method of impregnating an intraluminal prosthesis with a pharmacological agent, comprising:

immersing an intraluminal prosthesis in a mixture of a carrier fluid and a pharmacological agent, wherein the intraluminal prosthesis comprises non-layered polymeric material;

pressurizing the mixture of carrier fluid and pharmacological agent for a time sufficient to cause the carrier fluid and pharmacological agent to at least partially penetrate the non-layered polymeric material;

removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient, wherein the concentration gradient defines an elution profile of the pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject.

Appellants respectfully traverse the rejection of independent Claim 73 and claims dependent therefrom.

Igaki fails to teach or suggest removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient. In fact, the Final Action mailed February 12, 2008 (hereinafter the "Action") concedes that Igaki fails to teach a drug elutably trapped within polymeric material in a predetermined concentration gradient. (Action, Page 2). However, the Action states that "Igaki teaches that the pressure is gradually exhausted (i.e., removing pressure over a predetermined period of time) in a reaction chamber (i.e., under controlled conditions) and, thus, teaches all the steps as claimed." (Action, Page 3). On this basis, the Action then concludes that, "because the method of Igaki is so similar to the claimed steps, the two methods must necessarily achieve similar results." (Action, Page 3). Appellants respectfully disagree.

Igaki does not teach the same steps nor does it achieve similar results as those of the present invention. The Action cites to paragraph 0062 of Igaki for support for the assertion that Igaki "teaches all the steps as claimed" and therefore that "the two methods must necessarily achieve the same results." Action, page 3. Paragraph 0062 of Igaki recites:

Finally, a third valve 31 is opened to exhaust CO₂ within the reaction chamber 27 gradually to set the inside of the reaction chamber 27 open to atmosphere. The drug 26 is now **fully impregnated** in the stent 1 to complete the luminal stent according to the present invention.
(emphasis added)

Thus, a full and fair reading of paragraph 0062 of Igaki fails to show that the steps of Igaki result in a concentration gradient in the polymeric material as taught by the present invention but rather

describes a process in which a stent is "fully impregnated" with the drug." Clearly, one of ordinary skill in the art would not reasonably interpret this to mean the formation of a predetermined concentration gradient as taught by the present invention. Further, the data presented in Table 3 and in Figures 10 and 11 of Igaki support the conclusion that the methods of Igaki in fact teach that only a single concentration of drug is impregnated in the polymeric material of Igaki. For instance, in Table 3 each example presents only a single concentration of impregnated drug. Furthermore, commenting on the Examples, Igaki states "[i]t has been shown that the quantity of transilast impregnated in the stent depends on the pressure and temperature of the supercritical fluid CO₂, and that, in particular, if the temperature at which CO₂ is made into a supercritical fluid is high, the quantity of transilast impregnated is increased." (Igaki, Para. 98). No teaching or suggestion is made of controlling such variables as pressure or temperature so as to achieve a concentration gradient in the polymeric material but rather, Igaki describes only increasing or decreasing the absolute amount of drug impregnated in the stent. Thus, based on the data and text presented in Igaki et al., one of ordinary skill in the art would not reasonably conclude that Igaki et al. teaches a concentration gradient in the polymeric material.

The statements in the Action such as "must necessarily have some sort of concentration gradient within the polymeric material in the method of Igaki" and the "concentration gradient, therefore must necessarily define an elution profile of the pharmacological agent as required in the claims," appear to be based on a conclusion that Igaki inherently teaches removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient as is taught by the present invention. Appellants respectfully submit that, as stated in the M.P.E.P., a reference cannot be relied upon for allegedly inherent teachings to support a rejection under 35 U.S.C. §103. Specifically, it is stated in § 2141.02 (V) of the M.P.E.P., with a citation to *In re Rijckaert*, that "[o]bviousness cannot be predicated on what is not known at the time an invention was made, even if the inherency of a certain feature is later established." (28 USPQ2d 1955 (Fed. Cir. 1993)).

Contrary to the assertion in the Advisory Action mailed April 25, 2008, *In re Rijckaert* does, in fact, address the issue of the use of inherency in a 35 U.S.C. §103 rejection. In that case,

the court reversed the Board of Patent Appeals and Interferences (BPAI) decision to uphold an obviousness rejection that was based on a conclusion that the relationship between time compression/expansion and the three particular variables was inherent in the prior art (28 USPQ2d at 1957). The court agreed with the appellant that "the examiner's assumptions do not constitute the disclosure of prior art" and stated that "[t]he mere fact that a certain thing may result from a given set of circumstances is not sufficient [to establish inherency]." (*Id.*) The court then citing to *In re Spormann* stated "[t]hat which may be inherent is not necessarily known. Obviousness cannot be predicated on what is unknown." (*Id.*) Thus, the use of the Igaki reference for its allegedly inherent teachings as a necessary basis for this obviousness rejection renders the rejection improper and Appellants request that it be withdrawn for at least this reason.

Even if one assumes that using an allegedly inherent teaching as basis for an obviousness rejection is proper (which Appellants do not concede), the present rejection still fails because no extrinsic evidence is provided in the Action to make it clear that the missing descriptive matter is necessarily present as is required for such an argument. The legal standard for inherency is set forth in section 2112 in the MPEP and states that "[t]o establish inherency, the extrinsic evidence must make clear that the missing descriptive matter is necessarily present in the thing described in the reference, and that it would be so recognized by persons of ordinary skill. Inherency, however, may not be established by probabilities or possibilities. The mere fact that a certain thing may result from a given set of circumstances is not sufficient." (*In re Robertson* 169 F.3d 743, 745, 49 USPQ2d 1949, 1950-51 (Fed. Cir. 1999). (Emphasis added.) The MPEP also cites *Ex parte Levy* as stating that "[i]n relying upon the theory of inherency, the examiner must provide a basis in fact and/or technical reasoning to reasonably support the determination that the allegedly inherent characteristic necessarily flows from the teachings of the applied prior art." (17 USPQ2d 1461, 1464 (Bd. Pat. App. & Inter. 1990)). It is clear from these cases that in order for the standard for inherency to be met, the allegedly inherent characteristic must necessarily flow from the teachings of the cited art and such a determination must be supported by fact or technical reasoning.

In the present case, there is no teaching or suggestion in the Igaki reference that the pressure was necessarily removed over a predetermined period of time and under controlled

conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient as is taught by the present invention. This is mere speculation on the Examiner's part and there is no fact or technical reasoning provided in support of this statement, as required to demonstrate that the methods of the claimed invention are inherently disclosed in the Igaki reference. Thus, there is no evidence that the claimed methods necessarily flow from the teachings of the cited art and therefore the claimed methods of this invention cannot be obvious as inherent in the teachings of Igaki.

Furthermore, contrary to the assertions in the Action, the only method provided in Igaki for controlling the release time point and quantity of an impregnated drug is via the use of layers of biodegradable polymer material and not via a concentration gradient. Igaki specifically states "[b]y forming a further biodegradable polymer layer on the stent surface, it becomes possible to control the release rate of the drug into the blood, impregnated inside of the stent." (Igaki, Para, 0026). Igaki also states "[b]y providing the layer(s) of the drug-containing biodegradable polymer in this manner, one or more drugs may be impregnated in the stent, and it is possible to permit more strict control of the drug releasing time point or the quantity of the released drug, or different drugs can be released at the desired same time point." (Igaki, Para. 0072). In Paragraph 0067 of Igaki it is further stated "...[f]or delaying the release of the drug impregnated in the stent into the blood vessel, it is also possible to form the layer of the biodegradable polymer material, formed only of the biodegradable polymer...." (Igaki,) Accordingly, Igaki fails to teach or suggest a method of impregnating an interluminal prosthesis comprising removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and *the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient*, as taught by the present invention. In fact, Appellants submit that the teachings of Igaki of the use of layers for controlling the release time point and quantity of an impregnated drug teaches away from the use of concentration gradients.

Appellants note that the Action further states that a predetermined concentration gradient can be zero. (Action, Page 3). Appellants respectfully disagree. One of skill in the art would know that the term gradient is defined as "the rate of regular or graded ascent or descent" and

that a "concentration gradient" is defined as "a gradual change in the concentration of solutes in a solution as a function of distance through a solution" (see the dictionary definitions included herein in the Evidence Appendix). A concentration can be zero; however, a gradient by definition cannot be zero. Clearly one of ordinary skill in the art would not reasonably consider a concentration gradient to be zero.

Thus, one of ordinary skill in the art would not, upon reading Igaki, produce a stent comprising a non-layered polymeric material with a drug concentration gradient in the nonlayered polymeric material as taught in the present application. To the contrary, one skilled in the art, upon reading Igaki, would use various layers to accomplish what Appellants accomplish via a concentration gradient. The fact that Igaki utilizes layers to control the time of drug release and the quantity of drug release illustrates that Igaki does not utilize or appreciate utilizing a drug concentration gradient for accomplishing time of drug release and/or quantity of drug release.

In view of the foregoing, Appellants respectfully submit that the rejection under 35 U.S.C. §103(a) of independent Claim 73 and claims dependent thereon, Claims 74, 76, 80-84, and 86, over Igaki is overcome and request that this rejection be withdrawn.

B. Dependent Claim 86.

Claim 86 depends from independent Claim 73 and recites wherein the non-layered polymeric material is a coating on a portion of the intraluminal prosthesis.

As discussed above, Igaki fails to teach or suggest all of the recitations of independent Claim 73. With regard to dependent Claim 86, the Action states that the polymeric material can be formed only on the surface. The Action provides no support for this assertion and none can be found in Igaki. At most, Igaki discusses a further layer of polymeric material on the surface of the impregnated stent in order to control the rate of release of the drug impregnated in the stent. (Igaki, *see, e.g.*, Abstract, para. 0017-0027). Thus, there is no teaching or suggestion in Igaki that the polymeric material is a coating on a portion of the intraluminal prosthesis as recited in Claim 86 of the present invention.

Therefore, dependent Claim 86 is independently patentable over Igaki and Appellants respectfully request withdrawal of these rejections.

In sum, Igaki fails to teach or suggest all of the recitations of independent Claim 73 and dependent claims 74, 76, 80-84 and 86, and Appellants respectfully request withdrawal of this rejection.

III. Claims 73, 74, 76-78, 80-82, 86, 88, 89, 91-93 and 98 are patentable over European Patent No. EP 0405284 to Greiner.

In addition, Claims 73, 74, 76-78, 80-82, 86, 88, 89, 91-93 and 98 stand rejected under 35 U.S.C. §103(a) as allegedly unpatentable over European Patent No. EP 0405284 to Greiner (hereinafter "Greiner"). Appellants respectfully disagree and request reversal of this rejection.

A. Independent Claim 73.

The recitations of Appellants' independent Claim 73 are set forth above in section II. Appellants respectfully traverse the rejection of independent Claim 73 and claims dependent therefrom.

Greiner describes a method of impregnating a catheter, made of polymeric material, with a pharmaceutical. The catheter is immersed into a saturated solution of a pharmaceutical in a solvent. The saturated solution serves as a swelling agent and swells the polymeric material of the catheter. The catheter is contacted with the swelling agent at or near supercritical pressure and temperature of the solvent. The pressure is then reduced from supercritical pressure to release the solvent from the catheter, thereby leaving the pharmaceutical impregnated within the catheter. Greiner fails to teach or suggest removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient.

The Action states that the polymeric material of Greiner would "necessarily have a concentration gradient of the pharmacological agent for substantially the same reasons as discussed..." (for Igaki). Action, Page 4. Further, the Action states once again that a concentration gradient can be zero. *Id.* Appellants respectfully disagree.

Similar to Igaki, Greiner also fails to teach or suggest an intraluminal prosthesis wherein *a pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient* as taught in the present application. In fact, the

catheter of the Greiner invention is discussed only in terms of being impregnated with specific concentrations of a drug. Thus, Greiner states that "[t]he catheter is found to contain 25% benzocaine by weight...." (Greiner, Col. 5, Example 1). In Example 2, Greiner states that the catheter "is found to contain 42% benzocaine." Thus, Greiner provides no teaching or suggestion of a concentration **gradient** within a non-layered polymeric material as is taught by the present invention.

Further, contrary to the allegations in the Action, Greiner does not describe removing the pressure over a predetermined time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient as recited in Claim 73 of the present application. The Action cites to column 4, lines 2-6 of Greiner wherein it states "[a]fter contacting, the volatile swelling agent is separated from the catheter, leaving the pharmaceutical behind. Because of the volatility of the swelling agents employed, separation is easily accomplished by lowering the pressure." Action, Page 5. The Action then contends that "[t]here must be some control of how fast the rate of pressure changes" (*Id.*) but does not provide any evidence for such a supposition.

As discussed above, in section IIA, a reference cannot be relied upon for allegedly inherent teachings to support a rejection under 35 U.S.C. §103. Specifically, it is stated in § 2141.02 (V) of the M.P.E.P., with a citation to *In re Rijckaert*, that "[o]bviousness cannot be predicated on what is not known at the time an invention was made, even if the inherency of a certain feature is later established." (28 USPQ2d 1955 (Fed. Cir. 1993)). Furthermore, even if one assumes that using an allegedly inherent teaching as basis for an obviousness rejection is proper (which Appellants do not concede), the present rejection still fails because, as pointed out above, no extrinsic evidence is provided in the Action to make it clear that the missing descriptive matter is necessarily present in Greiner as is required for such an argument.

As stated in the Manual for Patent Examining Procedure § 2112 (V) when an examiner relies upon a theory of inherency, "the examiner must provide a basis in fact and/or technical reasoning to reasonably support the determination that the allegedly inherent characteristic necessarily flows from the teachings of the applied prior art." *Ex parte Levy*, 17 USPQ2d 1461, 1464 (Bd. Pat. App. & Int. 1990). Inherency "may not be established by probabilities or

possibilities. The mere fact that a certain thing may result from a given set of circumstances is not sufficient." Ex parte Skinner, 2USPQ2d 1788, 1789 (Bd. Pat. App. & Int. 1986). Also, the examiner has the initial burden of providing such evidence or technical reasoning. See In re Spada, 911 F.2d 705, 708, 15 USPQ2d. 1655, 1657 (Fed. Cir. 1990); In re King, 801 F.2d 1324, 1327, 231 USPQ 136, 138-139 (Fed. Cir. 1986). In the present rejection, no such basis in fact and/or technical reasoning has been provided to reasonably support the allegation that Greiner inherently describes removing the pressure over a predetermined time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient as recited in Claim 73 of the present application.

In view of the foregoing, Appellants respectfully submit that the rejection under 35 U.S.C. §103(a) of independent Claim 73 and claims dependent thereon, Claims 74, 76-78, 80-82, and 86, over Greiner are overcome and request that these rejections be withdrawn.

B. Dependent Claim 86

Claim 86 depends from independent Claim 73 and recites wherein the non-layered polymeric material is a coating on a portion of the intraluminal prosthesis.

As discussed above, Greiner fails to teach or suggest all of the recitations of independent Claim 73. With regard to dependent Claim 86, the Action states that the polymeric material can be formed only on the surface. The Action provides no support for this assertion and none can be found in Greiner. At most, Greiner describes a method of impregnating a catheter, made of polymeric material, with a pharmaceutical. The catheter is immersed into a saturated solution of a pharmaceutical in a solvent. The saturated solution serves as a swelling agent and swells the polymeric material of the catheter. The catheter is contacted with the swelling agent at or near supercritical pressure and temperature of the solvent. The pressure is then reduced from supercritical pressure to release the solvent from the catheter, thereby leaving the pharmaceutical impregnated within the catheter. Thus, there is no teaching or suggestion in Greiner that the polymeric material is a coating on a portion of the intraluminal prosthesis as recited in Claim 86 of the present invention.

Accordingly, dependent Claim 86 is independently patentable over Greiner and Appellants respectfully request withdrawal of these rejections.

C. Independent Claim 88

Appellants' independent Claim 88 recites a method of impregnating an intraluminal prosthesis with a pharmacological agent, comprising:

- immersing an intraluminal prosthesis in a mixture of a carrier fluid and a pharmacological agent, wherein the intraluminal prosthesis comprises non-layered polymeric material;

- placing the intraluminal prosthesis within a pressure vessel;

- pressurizing the interior of the pressure vessel with an inert gas to a predetermined pressure, wherein the inert gas is selected from the group consisting of helium, nitrogen, and argon;

- supplying a mixture of a carrier fluid and a pharmacological agent into the pressure vessel;

- exposing the non-layered polymeric material and the mixture of carrier fluid and pharmacological agent in the pressure vessel for a time such that the carrier fluid and pharmacological agent at least partially penetrate the non-layered polymeric material; and

- releasing the pressure in the pressure vessel over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient, wherein the concentration gradient defines an elution profile of the pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject.

As discussed above, Greiner fails to teach or suggest removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration as taught by the presently claimed invention. Thus, for at least the same reason that Greiner fails to teach or suggest all of the recitations of independent Claim 73, as discussed above, Greiner fails to teach or suggest all of the recitations of independent Claim 88 so as to provide a motivation or reasonable expectation of success that would lead one of ordinary skill in the art, at the time this invention was made, to carry out the methods of the claimed invention. As such, Appellants respectfully assert that the 35 U.S.C. §103 rejection of independent Claim 88 and claims depending therefrom, Claims 89, 91-93 and 98, is overcome, and thus respectfully request its withdrawal.

D. Dependent Claim 98

Claim 98 depends from independent Claim 88 and recites wherein the non-layered polymeric material is a coating on a portion of the intraluminal prosthesis.

As discussed above, Greiner fails to teach or suggest all of the recitations of independent Claim 88. With regard to dependent Claim 98, the Action states that the polymeric material can be formed only on the surface. The Action provides no support for this assertion and none can be found in Greiner. Greiner describes a method of impregnating a catheter, made of polymeric material, with a pharmaceutical. The catheter is immersed into a saturated solution of a pharmaceutical in a solvent. The saturated solution serves as a swelling agent and swells the polymeric material of the catheter. The catheter is contacted with the swelling agent at or near supercritical pressure and temperature of the solvent. The pressure is then reduced from supercritical pressure to release the solvent from the catheter, thereby leaving the pharmaceutical impregnated within the catheter. Thus, there is no teaching or suggestion that the polymeric material is a coating on a portion of the intraluminal prosthesis as recited in Claim 98 of the present invention.

Accordingly, dependent Claim 98 is independently patentable over Greiner and Appellants respectfully request withdrawal of the rejection of Claim 98 over Greiner.

In sum, Greiner fails to teach or suggest all of the recitations of independent Claim 73 and dependent claims 74, 76, 80-84, 86, 88, 89, 91-93 and 98 and Appellants respectfully request withdrawal of this rejection.

IV. Claims 75, 99-101 and 104 are patentable over U.S. Patent Publication No. 2003/0104030 to Igaki et al. in view of U.S. Patent No. 6,251,136 to Guruwaiya.

Claims 75, 99-101 and 104 stand rejected under 35 U.S.C. §103(a) as allegedly unpatentable over Igaki in view of U.S. Patent No. 6,251,136 to Guruwaiya (hereinafter "Guruwaiya"). Appellants respectfully disagree and request reversal of this rejection.

A. Dependent Claim 75

Claim 75 depends from independent Claim 73 and recites the method of Claim 73, further comprising masking one or more portions of the intraluminal prosthesis with a protective layer of material prior to immersing the intraluminal prosthesis in a mixture of a carrier fluid and a pharmacological agent, wherein the protective

layer of material is configured such that the mixture of carrier fluid and pharmacological agent at least partially penetrate only unmasked portions of the non-layered polymeric material during the pressurizing step..

As discussed above, Igaki fails to teach or suggest all of the recitations of independent Claim 73 (See section IIA). Further, Guruwaiya fails to remedy the deficiencies of Igaki. Guruwaiya simply discusses the use of known masking techniques. Therefore, Igaki and Guruwaiya, alone or in any combination, fail to teach or suggest all of the recitations of independent Claim 73 and dependent claim 75 so as to provide a motivation or reasonable expectation of success that would lead one of ordinary skill in the art, at the time this invention was made, to carry out the methods of the claimed invention. Accordingly, Appellants respectfully request reversal of this rejection.

B. Independent Claim 99 and dependent Claims 100, 101 and 104.

Appellants' independent Claim 99 recites a method of impregnating an intraluminal prosthesis with a pharmacological agent, comprising:

- masking portions of an intraluminal prosthesis with a protective layer of material such that the intraluminal prosthesis has first and second unmasked portions, wherein the intraluminal prosthesis comprises non-layered polymeric material;

- immersing the intraluminal prosthesis in a mixture of a carrier fluid and first and second pharmacological agents;

- pressurizing the mixture of carrier fluid and pharmacological agents for a time sufficient to cause the carrier fluid and the first pharmacological agent to at least partially penetrate the first unmasked portion and to cause the carrier fluid and the second pharmacological agent to at least partially penetrate the second unmasked portion; and

- removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and such that an amount of the first pharmacological agent remains elutably trapped within the first unmasked portion in a predetermined concentration gradient and an amount of the second pharmacological agent remains elutably trapped within the second unmasked portion in a predetermined concentration gradient, wherein each concentration gradient defines an elution profile of a respective pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject.

As discussed above (See section IIA), Igaki fails to teach or suggest removing the pressure over a predetermined period of time and under controlled conditions such that the

carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient. In fact, the text and data of Igaki support the conclusion that the methods of Igaki result in a polymeric material comprising a single concentration of drug. (See, Igaki, for example, at least, Para. 0026, 0072, 0067, Table 3, Figures 10 and 11). The only method provided by Igaki for controlling the release time point and quantity of an impregnated drug is not via a concentration gradient, but rather via the use of layers of biodegradable polymer material. Igaki specifically states "[b]y providing the layer(s) of the drug-containing biodegradable polymer in this manner, one or more drugs may be impregnated in the stent, and it is possible to permit more strict control of the drug releasing time point or the quantity of the released drug, or different drugs can be released at the desired same time point." (Igaki, Para. 0072). Igaki also describes retarding the rate of release of a drug impregnated in a stent via the use of a layer of biodegradable polymer material not containing a drug. (Igaki, Para. 0073).

One skilled in the art would not, upon reading Igaki, produce a non-layered stent with a drug concentration gradient in the stent material. To the contrary, one skilled in the art, upon reading Igaki, would use various layers to accomplish what Appellants accomplish via a concentration gradient. The fact that Igaki utilizes layers to control the time of drug release and the quantity of drug release illustrates that Igaki does not utilize or appreciate utilizing a drug concentration gradient for accomplishing time of drug release and/or quantity of drug release.

The secondary reference, Guruwaiya, fails to remedy the deficiencies of Igaki. The Action states that Guruwaiya teaches a method of coating a pharmacological agent on a stent wherein certain portions of the stent are masked during the coating process. (Action, Page 3). The Action then concludes that it would have been obvious to have masked certain portions of the stent of Igaki. (Action, Page 3). Appellants respectfully disagree.

Appellants submit that to the extent that the combinations of Igaki and Guruwaiya teach masking the stent of Igaki, such a combination fails to teach or suggest the following recitations of independent Claim 99:

masking portions of an intraluminal prosthesis with a protective layer of material such that the intraluminal prosthesis has first and second unmasked portions, wherein the intraluminal prosthesis comprises non-layered polymeric material;... and

removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and such that an amount of the first pharmacological agent remains elutably trapped within the first unmasked portion in a predetermined concentration gradient and an amount of the second pharmacological agent remains elutably trapped within the second unmasked portion in a predetermined concentration gradient, wherein each concentration gradient defines an elution profile of a respective pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject.

Accordingly, Igaki and Guruwaiya, alone or in combination, fail to teach or suggest all of the recitations of independent Claim 99 so as to provide a motivation or reasonable expectation of success that would lead one of ordinary skill in the art, at the time this invention was made, to carry out the methods of the claimed invention. Thus, appellants respectfully assert that the rejection under 35 U.S.C. §103 of independent Claim 99 and Claims 100, 101 and 104, depending therefrom, over Igaki in view of Guruwaiya is overcome.

In view of the foregoing, Appellants respectfully submit that the rejection under 35 U.S.C. §103(a) of Claims 75, 99-101 and 104 over Igaki in view of Guruwaiya is overcome and request that these rejections be withdrawn.

V. Claims 75, 90 and 99-104 are patentable over European Patent No. EP 0405284 to Greiner in view of U.S. Patent No. 6,251,136 to Guruwaiya.

Claims 75, 90 and 99-104 stand rejected under 35 U.S.C. §103(a) as allegedly unpatentable over Greiner in view of Guruwaiya. Appellants respectfully disagree and request reversal of this rejection.

A. Dependent Claim 75

Appellants' dependent Claim 75 depends from independent Claim 73 and recites the method of Claim 73 as set forth above, in section IV(A).

As discussed above, Greiner fails to teach or suggest all of the recitations of independent Claim 73 (See section IIIA). Further, Guruwaiya fails to remedy the deficiencies of Greiner. Guruwaiya simply discusses the use of known masking techniques. Therefore, Greiner and Guruwaiya, alone or in any combination, fail to teach or suggest all of the recitations of independent Claim 73 and dependent claim 75 so as to provide a motivation or reasonable

expectation of success that would lead one of ordinary skill in the art, at the time this invention was made, to carry out the method of Claim 75. Accordingly, Appellants respectfully request reversal of this rejection.

B. Dependent Claim 90

Appellants' dependent Claim 90 depends from independent Claim 88 and recites the method of Claim 88,

further comprising masking one or more portions of the intraluminal prosthesis with a protective layer of material prior to immersing the intraluminal prosthesis in a mixture of a carrier fluid and a pharmacological agent, wherein the protective layer of material is configured such that the mixture of carrier fluid and pharmacological agent at least partially penetrate only unmasked portions of the non-layered polymeric material during the pressurizing step.

As discussed above, Greiner fails to teach or suggest all of the recitations of independent Claim 88 (See section IIIB). Further, Guruwaiya fails to remedy the deficiencies of Greiner. Guruwaiya simply discusses the use of known masking techniques. Therefore, Greiner and Guruwaiya, alone or in any combination, fail to teach or suggest all of the recitations of independent Claim 88 and dependent claim 90 so as to provide a motivation or reasonable expectation of success that would lead one of ordinary skill in the art, at the time this invention was made, to carry out the methods of Claim 90. Accordingly, Appellants respectfully request reversal of this rejection.

C. Independent Claim 99 and dependent Claims 100-104.

Appellants' independent Claim 99 is set forth above, in section IV(B).

As discussed above (See section IIIA), Greiner fails to teach or suggest a drug elutably trapped within non-layered polymeric material in a predetermined concentration gradient as taught by the presently claimed invention. For at least the same reasons that Greiner fails to teach or suggest all of the recitations of independent Claims 73 and 88, as discussed above, Greiner also fails to teach or suggest all of the recitations of independent Claim 99.

The secondary reference, Guruwaiya, fails to remedy the deficiencies of primary reference, Greiner. The Action states that Guruwaiya teaches a method of coating a pharmacological agent on a stent wherein certain portions of the stent are masked during the coating process. (Action, Page 3). The Action then concludes that it would have been obvious to

have masked certain portions of the stent of Igaki. (Action, Page 3). The Action further states that Claim 99 is rejected under 35 U.S.C. 103(a) as being unpatentable over Greiner in view of Guruwaiya, for substantially the same reasons as discussed for Igaki in view of Guruwaiya. Appellants respectfully disagree.

Appellants submit that to the extent that the combination of Greiner and Guruwaiya teach masking the stent of Greiner, such a combination fails to teach or suggest the following recitations of independent Claim 99:

masking portions of an intraluminal prosthesis with a protective layer of material such that the intraluminal prosthesis has first and second unmasked portions, wherein the intraluminal prosthesis comprises non-layered polymeric material;... and

removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and such that an amount of the first pharmacological agent remains elutably trapped within the first unmasked portion in a predetermined concentration gradient and an amount of the second pharmacological agent remains elutably trapped within the second unmasked portion in a predetermined concentration gradient, wherein each concentration gradient defines an elution profile of a respective pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject.

Accordingly, Greiner and Guruwaiya, alone or in combination, fail to teach or suggest all of the recitations of independent Claim 99 so as to provide a motivation or reasonable expectation of success that would lead one of ordinary skill in the art, at the time this invention was made, to carry out the methods of the claimed invention. Thus, appellants respectfully assert that the rejection under 35 U.S.C. §103 of independent Claim 99, and Claims 100, 101 and 104 depending therefrom, over Greiner in view of Guruwaiya is overcome

In view of the foregoing, Appellants respectfully submit that the rejection under 35 U.S.C. §103(a) of Claims 75, 90, and 99-104 over Greiner in view of Guruwaiya is overcome and request that these rejections be withdrawn.

VI. Claim 79 is patentable over U.S. Patent Publication No. 2003/0104030 to Igaki et al. in view of U.S. Patent No. 6,670,398 to Edwards.

Claim 79 stands rejected under 35 U.S.C. §103(a) as allegedly unpatentable over Igaki in view of U.S. Patent No. 6,670,398 to Edwards (hereinafter "Edwards"). Appellants respectfully request reversal of this rejection.

Claim 79 depends from independent claim 73 and recites that the pharmacological agent comprises everolimus.

As discussed above, Igaki fails to teach or suggest all of the recitations of independent Claim 73 (See section IIA). Further, Edwards fails to remedy the deficiencies of Igaki. Edwards simply describes everolimus as a therapeutic drug that can be used to suppress a transplant recipient's immune response. Thus, the cited references, alone or in combination, fail to teach or suggest all of the recitations of the present invention or provide any reasonable expectation of success of achieving the present invention with such a combination. Accordingly, Appellants respectfully submit that the rejection of Claim 79 under 35 U.S.C. §103(a) over Igaki in view of Edwards is overcome and respectfully request its withdrawal.

VII. Claims 79 and 95 are patentable over European Patent No. EP 0405284 to Greiner in view of U.S. Patent No. 6,670,398 to Edwards.

Claims 79 and 95 stand rejected under 35 U.S.C. §103(a) as allegedly unpatentable over Greiner in view of Edwards. Appellants respectfully request reversal of these rejections.

Claim 79 depends from independent claim 73 and recites that the pharmacological agent comprises everolimus. Claim 95 depends from independent Claim 88 and recites that the pharmacological agent comprises everolimus.

As discussed above, Greiner fails to teach or suggest all of the recitations of independent Claim 73 and independent Claim 88 (See sections IIIA and IIIC). Further, Edwards fails to remedy the deficiencies of Greiner. Edwards simply describes everolimus as a therapeutic drug that can be used to suppress a transplant recipient's immune response. Thus, the cited references, alone or in combination, fail to teach or suggest all of the recitations of the present invention or provide any reasonable expectation of success of achieving the present invention with such a

combination. Accordingly, Appellants respectfully submit that the rejection of Claims 79 and 95 under 35 U.S.C. §103(a) over Greiner in view of Edwards is overcome and respectfully request its withdrawal.

VIII. Claims 85 and 97 are patentable over European Patent No. EP 0405284 to Greiner in view of PCT Publication WO 01/87368 to Mehta.

Claims 85 and 97 stand rejected under 35 U.S.C. §103(a) as allegedly unpatentable over Greiner in view of PCT Publication WO 01/87368 to Mehta (hereinafter "Mehta"). Appellants respectfully request reversal of this rejection.

Claim 85 depends from independent Claim 73 and recites the method wherein the non-layered polymeric material is non-erodible. Claim 97 depends from independent Claim 88 and recites the method wherein the non-layered polymeric material is non-erodible.

As discussed above, Greiner fails to teach or even suggest all of the recitations of Claims 73 and 88 (See sections IIIA and IIIC). Further, Mehta fails to remedy the deficiencies of Greiner. Mehta simply describes deposition of a coating by altering the temperature and pressure of a SCF in which a drug or polymer to be coated is dissolved. Thus, the cited references, alone or in combination, fail to teach or suggest all of the recitations of the present invention or provide any reasonable expectation of success of achieving the present invention with such a combination. Accordingly, Appellants respectfully submit that the rejection of Claims 85 and 97 under 35 U.S.C. §103(a) over Greiner in view of Mehta is overcome and respectfully request its withdrawal.

IX. Claim 87 is patentable over U.S. Patent Publication No. 2003/0104030 to Igaki et al. in view of U.S. Patent No. 6,299,604 to Ragheb.

Claim 87 stands rejected under 35 U.S.C. §103(a) as allegedly unpatentable over Igaki in view of U.S. Patent No. 6,299,604 to Ragheb (hereinafter "Ragheb").

Claim 87 depends from independent Claim 73 and recites the method further comprising immersing the intraluminal prosthesis in a mixture of a carrier fluid and radiopaque material; and pressurizing the mixture of carrier fluid and radiopaque material for a time sufficient to cause the

carrier fluid and radiopaque material to at least partially penetrate the non-layered polymeric material.

As discussed above, Igaki fails to teach or even suggest all of the recitations of independent Claim 73 (See section IIA). Further, Ragheb fails to remedy the deficiencies of Igaki. Ragheb simply describes radiopaque agents as alternative bioactive materials that can be used in the vascular system. Thus, the cited references, alone or in combination, fail to teach or suggest all of the recitations of the present invention or provide any reasonable expectation of success of achieving the present invention with such a combination. Accordingly, Appellants respectfully submit that the rejection of Claim 87 under 35 U.S.C. §103(a) over Igaki in view of Ragheb is overcome and respectfully request its withdrawal.

X. Claim 87 is patentable over European Patent No. EP 0405284 to Greiner in view of U.S. Patent No. 6,299,604 to Ragheb.

Claim 87 stands rejected under 35 U.S.C. §103(a) as allegedly unpatentable over Greiner in view of Ragheb. Appellants respectfully request reversal of this rejection.

Claim 87 depends from independent Claim 73 and recites the method further comprising immersing the intraluminal prosthesis in a mixture of a carrier fluid and radiopaque material; and pressurizing the mixture of carrier fluid and radiopaque material for a time sufficient to cause the carrier fluid and radiopaque material to at least partially penetrate the non-layered polymeric material.

As discussed above, Greiner fail to teach or even suggest all of the recitations of independent Claim 73 (See section IIIA). Further, Ragheb fails to remedy the deficiencies of Greiner. Ragheb simply describes radiopaque agents as alternative bioactive materials that can be used in the vascular system. Thus, the cited references, alone or in combination, fail to teach or suggest all of the recitations of the present invention or provide any reasonable expectation of success of achieving the present invention with such a combination. Accordingly, Appellants respectfully submit that the rejections of Claim 87 under 35 U.S.C. §103(a) over Greiner in view of Ragheb is overcome and respectfully requests its withdrawal.

XI. Claims 81, 83, 84, 86, 93, 94, 96 and 98 are patentable over European Patent No. EP 0405284 to Greiner in view of U.S. Patent Publication No. 2003/0104030 to Igaki et al.

Claims 81, 83, 84, 86, 93, 94, 96 and 98 stand rejected under 35 U.S.C. §103(a) as allegedly unpatentable over Greiner in view of Igaki. Appellants respectfully request reversal of these rejections.

Dependent Claim 81 recites the methods of Claims 73, 76, and 80, wherein the carbon dioxide contains one or more of a co-solvent, a surfactant, and a co-surfactant. Dependent Claim 83 recites the methods of Claims 73, 76, 80 and 81, wherein the co-solvent is selected from the group consisting of ethanol and methanol. Dependent Claim 84 recites the method of Claim 73, wherein the intraluminal prosthesis is a stent. Dependent Claim 86 recites the method of Claim 73, wherein the non-layered polymeric material is a coating on a portion of the intraluminal prosthesis. Dependent Claim 93 recites the methods of Claims 88 and 91, wherein the carbon dioxide contains one or more of a co-solvent, a surfactant, and a co-surfactant. Dependent Claim 94 recites the methods of Claims 88, 91 and 93, wherein the co-solvent is selected from the group consisting of ethanol and methanol. Dependent Claim 96 recites the method of Claim 88, wherein the intraluminal prosthesis is a stent. Dependent Claim 98 recites the method of Claim 88, wherein the non-layered polymeric material is a coating on a portion of the intraluminal prosthesis.

As discussed above, Igaki and Greiner fail to teach or suggest all of the recitations of independent Claims 73 and 88 (See sections IIA, IIIA and IIIC). Thus, Igaki and Greiner, alone or in combination, fail to teach or suggest all of the recitations of the present invention or provide any reasonable expectation of success with such a combination. Accordingly, Appellants respectfully submit that the rejections of dependent Claims 81, 83, 84, 86, 93, 94, 96 and 98 under 35 U.S.C. §103(a) over Igaki in view of Ragheb and over Greiner in view of Ragheb are overcome and respectfully request their withdrawal.

A. Dependent Claim 86

Further, with regard to dependent Claim 86 (dependent from Claim 73), the Action states that the polymeric material can be formed only on the surface. The Action provides no support for this assertion and none can be found in either Igaki and Greiner. At most, Igaki discusses a further layer of polymeric material on the surface of the impregnated stent in order to control the

rate of release of the drug impregnated in the stent. (Igaki, *see, e.g.*, Abstract, para. 0017-0027). Greiner describes a method of impregnating a catheter, made of polymeric material, with a pharmaceutical. The catheter is immersed into a saturated solution of a pharmaceutical in a solvent. The saturated solution serves as a swelling agent and swells the polymeric material of the catheter. The catheter is contacted with the swelling agent at or near supercritical pressure and temperature of the solvent. The pressure is then reduced from supercritical pressure to release the solvent from the catheter, thereby leaving the pharmaceutical impregnated within the catheter. Thus, there is no teaching or suggestion in either Igaki or Greiner that the polymeric material is a coating on a portion of the intraluminal prosthesis as recited in Claim 86 of the present invention.

Accordingly, dependent Claim 86 is independently patentable over Igaki and Greiner, alone or in combination, and Appellants respectfully request withdrawal of this rejection.

B. Dependent Claim 98

Furthermore, with regard to dependent Claim 98 (dependent from Claim 88), the Action states that the polymeric material can be formed only on the surface. The Action provides no support for this assertion and none can be found in Greiner (or Igaki). Greiner describes a method of impregnating a catheter, made of polymeric material, with a pharmaceutical. The catheter is immersed into a saturated solution of a pharmaceutical in a solvent. The saturated solution serves as a swelling agent and swells the polymeric material of the catheter. The catheter is contacted with the swelling agent at or near supercritical pressure and temperature of the solvent. The pressure is then reduced from supercritical pressure to release the solvent from the catheter, thereby leaving the pharmaceutical impregnated within the catheter. Thus, there is no teaching or suggestion that the polymeric material is a coating on a portion of the intraluminal prosthesis as recited in Claim 98 of the present invention.

Accordingly, dependent Claim 98 is independently patentable over Greiner and Igaki and Appellants respectfully request withdrawal of this rejection.

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Conclusion

In light of the entire record and the above discussion, Appellants respectfully submit that each of the pending claims is patentable over the cited references and therefore request reversal of the rejections of Claims 73-104 and that this case be passed to issuance.

Respectfully submitted,



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CERTIFICATION OF ELECTRONIC TRANSMISSION UNDER 37 CFR § 1.8

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October 1, 2008.



Anthony DeRosa

Claims Appendix

1-72 (Cancelled)

73. (Previously Presented) A method of impregnating an intraluminal prosthesis with a pharmacological agent, comprising:

immersing an intraluminal prosthesis in a mixture of a carrier fluid and a pharmacological agent, wherein the intraluminal prosthesis comprises non-layered polymeric material;

pressurizing the mixture of carrier fluid and pharmacological agent for a time sufficient to cause the carrier fluid and pharmacological agent to at least partially penetrate the non-layered polymeric material;

removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient, wherein the concentration gradient defines an elution profile of the pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject.

74. (Previously Presented) The method of Claim 73, wherein the step of removing pressure is carried out under controlled conditions in which at least one parameter selected from the group consisting of temperature, rate of temperature change, pressure, rate of pressure change, carrier fluid quantity, and rate of carrier fluid quantity, is controlled in a predetermined pattern.

75. (Previously Presented) The method of Claim 73, further comprising masking one or more portions of the intraluminal prosthesis with a protective layer of material prior to immersing the intraluminal prosthesis in a mixture of a carrier fluid and a pharmacological agent, wherein the protective layer of material is configured such that the mixture of carrier fluid and

pharmacological agent at least partially penetrate only unmasked portions of the non-layered polymeric material during the pressurizing step.

76. (Previously Presented) The method of Claim 73, wherein the carrier fluid is carbon dioxide.

77. (Previously Presented) The method of Claim 73, wherein the carrier fluid is carbon dioxide and wherein the step of pressurizing the mixture of carrier fluid and pharmacological agent is performed using an inert second gas.

78. (Previously Presented) The method of Claim 77, wherein the second gas is selected from the group consisting of helium, nitrogen, and argon.

79. (Previously Presented) The method of Claim 73, wherein the pharmacological agent comprises everolimus.

80. (Previously Presented) The method of Claim 76, wherein the carbon dioxide is present in a supercritical state.

81. (Previously Presented) The method of Claim 80, wherein the carbon dioxide contains one or more of a co-solvent, a surfactant, and a co-surfactant.

82. (Previously Presented) The method of Claim 73, wherein the carrier fluid is configured to alter diffusion coefficients of the non-layered polymeric material.

83. (Previously Presented) The method of Claim 81, wherein the co-solvent is selected from the group consisting of ethanol and methanol.

84. (Previously Presented) The method of Claim 73, wherein the intraluminal prosthesis is a stent.

85. (Previously Presented) The method of Claim 73, wherein the non-layered polymeric material is non-erodible.

86. (Previously Presented) The method of Claim 73, wherein the non-layered polymeric material is a coating on a portion of the intraluminal prosthesis.

87. (Previously Presented) The method of Claim 73, further comprising:
immersing the intraluminal prosthesis in a mixture of a carrier fluid and radiopaque material; and
pressurizing the mixture of carrier fluid and radiopaque material for a time sufficient to cause the carrier fluid and radiopaque material to at least partially penetrate the non-layered polymeric material.

88. (Previously Presented) A method of impregnating an intraluminal prosthesis with a pharmacological agent, comprising:

immersing an intraluminal prosthesis in a mixture of a carrier fluid and a pharmacological agent, wherein the intraluminal prosthesis comprises non-layered polymeric material;

placing the intraluminal prosthesis within a pressure vessel;

pressurizing the interior of the pressure vessel with an inert gas to a predetermined pressure, wherein the inert gas is selected from the group consisting of helium, nitrogen, and argon;

supplying a mixture of a carrier fluid and a pharmacological agent into the pressure vessel;

exposing the non-layered polymeric material and the mixture of carrier fluid and pharmacological agent in the pressure vessel for a time such that the carrier fluid and pharmacological agent at least partially penetrate the non-layered polymeric material; and

releasing the pressure in the pressure vessel over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric

material and the pharmacological agent becomes elutably trapped within the non-layered polymeric material in a predetermined concentration gradient, wherein the concentration gradient defines an elution profile of the pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject.

89. (Previously Presented) The method of Claim 88, wherein the step of releasing pressure is carried out under controlled conditions in which at least one parameter selected from the group consisting of temperature, rate of temperature change, pressure, rate of pressure change, carrier fluid quantity, and rate of carrier fluid quantity, is controlled in a predetermined pattern.

90. (Previously Presented) The method of Claim 88, further comprising masking one or more portions of the intraluminal prosthesis with a protective layer of material prior to immersing the intraluminal prosthesis in a mixture of a carrier fluid and a pharmacological agent, wherein the protective layer of material is configured such that the mixture of carrier fluid and pharmacological agent at least partially penetrate only unmasked portions of the non-layered polymeric material during the pressurizing step.

91. (Previously Presented) The method of Claim 88, wherein the carrier fluid is carbon dioxide.

92. (Previously Presented) The method of Claim 91, wherein the carbon dioxide is in a supercritical state.

93. (Previously Presented) The method of Claim 91, wherein the carbon dioxide contains one or more of a co-solvent, a surfactant, and a co-surfactant.

94. (Previously Presented) The method of Claim 93, wherein the co-solvent is selected from the group consisting of ethanol and methanol.

95. (Previously Presented) The method of Claim 88, wherein the pharmacological agent is everolimus.

96. (Previously Presented) The method of Claim 88, wherein the intraluminal prosthesis is a stent.

97. (Previously Presented) The method of Claim 88, wherein the non-layered polymeric material is non-erodible.

98. (Previously Presented) The method of Claim 88, wherein the non-layered polymeric material is a coating on a portion of the intraluminal prosthesis.

99. (Previously Presented) A method of impregnating an intraluminal prosthesis with a pharmacological agent, comprising:

masking portions of an intraluminal prosthesis with a protective layer of material such that the intraluminal prosthesis has first and second unmasked portions, wherein the intraluminal prosthesis comprises non-layered polymeric material;

immersing the intraluminal prosthesis in a mixture of a carrier fluid and first and second pharmacological agents;

pressurizing the mixture of carrier fluid and pharmacological agents for a time sufficient to cause the carrier fluid and the first pharmacological agent to at least partially penetrate the first unmasked portion and to cause the carrier fluid and the second pharmacological agent to at least partially penetrate the second unmasked portion; and

removing the pressure over a predetermined period of time and under controlled conditions such that the carrier fluid diffuses out of the non-layered polymeric material and such that an amount of the first pharmacological agent remains elutably trapped within the first unmasked portion in a predetermined concentration gradient and an amount of the second pharmacological agent remains elutably trapped within the second unmasked portion in a predetermined concentration gradient, wherein each concentration gradient defines an elution

profile of a respective pharmacological agent from the non-layered polymeric material when the intraluminal prosthesis is deployed within a body of a subject.

100. (Previously Presented) The method of Claim 99, wherein the step of removing pressure is carried out under controlled conditions in which at least one parameter selected from the group consisting of temperature, rate of temperature change, pressure, rate of pressure change, carrier fluid quantity, and rate of carrier fluid quantity, is controlled in a predetermined pattern.

101. (Previously Presented) The method of Claim 99, wherein the carrier fluid is carbon dioxide.

102. (Previously Presented) The method of Claim 99, wherein the carrier fluid is carbon dioxide and wherein the step of pressurizing the mixture of carrier fluid and pharmacological agent is performed using an inert second gas.

103. (Previously Presented) The method of Claim 102, wherein the second gas is selected from the group consisting of helium, nitrogen, and argon.

104. (Previously Presented) The method of Claim 101, wherein the carbon dioxide is present in a supercritical state.

Evidence Appendix

Evidence listing:

Concentration definitions from:

- Biology-Online.org
- Merriam Webster Online Dictionary
- Online Medical Dictionary
- Answers.com

These documents were submitted concurrently with the response dated April 18, 2008.

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Related Proceedings Appendix

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